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**In the Claims:**

**Claim 1 (currently amended):** A method of fabricating a semiconductor device, having a reduced-oxygen copper-zinc alloy (Cu-Zn) thin film formed on a copper (Cu) surface by electroplating the Cu surface in a chemical solution, comprising the steps of:

providing forming a semiconductor substrate having a Cu surface on a semiconductor substrate;

providing a chemical solution;

electroplating the Cu surface in the chemical solution, thereby forming a Cu-Zn alloy thin film on the Cu surface, wherein the Cu-Zn alloy thin film completely covers the Cu surface;

rinsing the Cu-Zn alloy thin film in a solvent;

drying the Cu-Zn alloy thin film under a gaseous flow;

annealing the Cu-Zn alloy thin film formed on the Cu surface, thereby forming a reduced-oxygen Cu-Zn alloy thin film; and

completing formation of the semiconductor device.

**Claim 2 (original):** A method, as recited in Claim 1, wherein the chemical solution is nontoxic and aqueous, and wherein the chemical solution comprises:

at least one zinc (Zn) ion source for providing a plurality of Zn ions;

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at least one copper (Cu) ion source for providing a plurality of Cu ions;  
at least one complexing agent for complexing the plurality of Cu ions;  
at least one pH adjuster;  
at least one wetting agent for stabilizing the chemical solution, all being dissolved in a volume of deionized (DI) water.

**Claim 3 (currently amended):** A method, as recited in Claim 2,

wherein the at least one zinc (Zn) ion source comprises at least one zinc salt selected from a group consisting essentially of zinc acetate ((CH<sub>3</sub>CO<sub>2</sub>)<sub>2</sub>Zn), zinc bromide (ZnBr<sub>2</sub>), zinc carbonate hydroxide (ZnCO<sub>3</sub>·2Zn(OH)<sub>2</sub>), zinc dichloride (ZnCl<sub>2</sub>), zinc citrate ((O<sub>2</sub>CCH<sub>2</sub>C(OH)(CO<sub>2</sub>)CH<sub>2</sub>CO<sub>2</sub>)<sub>2</sub>Zn), zinc iodide (ZnI<sub>2</sub>), zinc L-lactate ((CH<sub>3</sub>CH(OH)CO<sub>2</sub>)<sub>2</sub>Zn), zinc nitrate (Zn(NO<sub>3</sub>)<sub>2</sub>), zinc stearate ((CH<sub>3</sub>(CH<sub>2</sub>)<sub>16</sub>CO<sub>2</sub>)<sub>2</sub>Zn), zinc sulfate (ZnSO<sub>4</sub>), zinc sulfide (ZnS), zinc sulfite (ZnSO<sub>3</sub>), and their hydrates.

**Claim 4 (currently amended):** A method, as recited in Claim 2,

wherein the at least one copper (Cu) ion source comprises at least one copper salt selected from a group consisting essentially of copper(I) acetate (CH<sub>3</sub>CO<sub>2</sub>Cu), copper(II) acetate ((CH<sub>3</sub>CO<sub>2</sub>)<sub>2</sub>Cu), copper(I) bromide (CuBr), copper (II) bromide (CuBr<sub>2</sub>), copper (II) hydroxide (Cu(OH)<sub>2</sub>), copper (II) hydroxide phosphate (Cu<sub>2</sub>(OH)PO<sub>4</sub>), copper(I) iodide (CuI), copper (II) nitrate ((CuNO<sub>3</sub>)<sub>2</sub>), copper(II) sulfate (CuSO<sub>4</sub>), copper(I) sulfide (Cu<sub>2</sub>S), copper(II) sulfide (CuS), copper (II) tartrate ((CH(OH)CO<sub>2</sub>)<sub>2</sub>Cu), and their

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hydrates.

**Claim 5 (original):** A method, as recited in Claim 1,

wherein said electroplating step comprises using an electroplating apparatus, and  
wherein said electroplating apparatus comprises:

- (a) a cathode-wafer;
- (b) an anode;
- (c) an electroplating vessel; and
- (d) a voltage source.

**Claim 6 (currently amended):** A method, as recited in Claim 5,

wherein the cathode-wafer comprises the Cu surface, and  
wherein the anode comprises at least one material selected from a group consisting  
essentially of copper (Cu), a copper-platinum alloy (Cu-Pt), titanium (Ti), platinum (Pt), a  
titanium-platinum alloy (Ti-Pt), an anodized copper-zinc alloy (Cu-Zn, i.e., brass), a  
platinized titanium (Pt/Ti), and a platinized copper-zinc (Pt/Cu-Zn, i.e., platinized brass).

**Claim 7 (canceled).**

**Claim 8 (currently amended):** A method, as recited in Claim 5,

wherein said electroplating comprises a plating condition selected from a group

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consisting essentially of a direct voltage in the range of approximately 1 V to approximately 4 V and a direct current in the range of approximately 0.01 A to approximately 0.2 A.

**Claim 9 (currently amended):** A method, as recited in Claim 5, wherein the Zn-doping in the reduced-oxygen Cu-Zn alloy thin film is controllable by varying at least one electroplating condition selected from a group consisting essentially of:

increasing the at least one zinc (Zn) ion source concentration, thereby slowly increasing said Zn-doping;

increasing the at least one copper (Cu) ion source concentration, thereby slowly decreasing said Zn-doping;

increasing the solution flow rate increases Zn-doping, thereby increasing the pH decreases cathodic efficiency with respect to Zn, and thereby decreasing said Zn-doping;

increasing the electroplating duration, thereby slowly decreasing said Zn-doping;

using a Cu anode, thereby decreasing said Zn-doping;

using a brass anode, thereby increasing said Zn-doping;

increasing the voltage, thereby increasing the Zn-doping; and

increasing the current, thereby increasing the Zn-doping.

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**Claim 10 (original):** A method, as recited in Claim 1,  
wherein the annealing step is performed in a temperature range of approximately  
150°C to approximately 450°C, and  
wherein the annealing step is performed for a duration range of approximately 0.5  
minutes to approximately 60 minutes.

**Claim 11 (currently amended):** A semiconductor device, having a reduced-oxygen copper-zinc alloy (Cu-Zn) thin film formed on a copper (Cu) surface by electroplating the Cu surface in a chemical solution, fabricated by a method comprising the steps of:

providing forming a semiconductor substrate having a Cu surface on a semiconductor substrate;  
providing a chemical solution;  
electroplating the Cu surface in the chemical solution, thereby forming a Cu-Zn alloy thin film on the Cu surface, wherein the Cu-Zn alloy thin film completely covers the Cu surface;  
rinsing the Cu-Zn alloy thin film in a solvent;  
drying the Cu-Zn alloy thin film under a gaseous flow;  
annealing the Cu-Zn alloy thin film formed on the Cu surface, thereby forming a reduced-oxygen Cu-Zn alloy thin film; and  
completing formation of the semiconductor device.

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**Claim 12 (original):** A device, as recited in Claim 11,  
wherein the chemical solution is nontoxic and aqueous, and  
wherein the chemical solution comprises:

at least one zinc (Zn) ion source for providing a plurality of Zn ions;  
at least one copper (Cu) ion source for providing a plurality of Cu ions;  
at least one complexing agent for complexing the plurality of Cu ions;  
at least one pH adjuster;  
at least one wetting agent for stabilizing the chemical solution, all being  
dissolved in a volume of deionized (DI) water.

**Claim 13 (currently amended):** A device, as recited in Claim 12,  
wherein the at least one zinc (Zn) ion source comprises at least one zinc salt  
selected from a group consisting essentially of zinc acetate ((CH<sub>3</sub>CO<sub>2</sub>)<sub>2</sub>Zn), zinc bromide  
(ZnBr<sub>2</sub>), zinc carbonate hydroxide (ZnCO<sub>3</sub>·2Zn(OH)<sub>2</sub>), zinc dichloride (ZnCl<sub>2</sub>), zinc  
citrate ((O<sub>2</sub>CCH<sub>2</sub>C(OH)(CO<sub>2</sub>)CH<sub>2</sub>CO<sub>2</sub>)<sub>2</sub>Zn<sub>3</sub>), zinc iodide (ZnI<sub>2</sub>), zinc L-lactate  
((CH<sub>3</sub>CH(OH)CO<sub>2</sub>)<sub>2</sub>Zn), zinc nitrate (Zn(NO<sub>3</sub>)<sub>2</sub>), zinc stearate ((CH<sub>3</sub>(CH<sub>2</sub>)<sub>16</sub>CO<sub>2</sub>)<sub>2</sub>Zn),  
zinc sulfate (ZnSO<sub>4</sub>), zinc sulfide (ZnS), zinc sulfite (ZnSO<sub>3</sub>), and their hydrates.

**Claim 14 (currently amended):** A device, as recited in Claim 12,  
wherein the at least one copper (Cu) ion source comprises at least one copper salt

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selected from a group consisting essentially of copper(I) acetate ( $\text{CH}_3\text{CO}_2\text{Cu}$ ), copper(II) acetate ( $(\text{CH}_3\text{CO}_2)_2\text{Cu}$ ), copper(I) bromide ( $\text{CuBr}$ ), copper (II) bromide ( $\text{CuBr}_2$ ), copper (II) hydroxide ( $\text{Cu}(\text{OH})_2$ ), copper (II) hydroxide phosphate ( $\text{Cu}_2(\text{OH})\text{PO}_4$ ), copper(I) iodide ( $\text{CuI}$ ), copper (II) nitrate ( $(\text{CuNO}_3)_2$ ), copper(II) sulfate ( $\text{CuSO}_4$ ), copper(I) sulfide ( $\text{Cu}_2\text{S}$ ), copper(II) sulfide ( $\text{CuS}$ ), copper (II) tartrate ( $(\text{CH}(\text{OH})\text{CO}_2)_2\text{Cu}$ ), and their hydrates.

**Claim 15 (original):** A device, as recited in Claim 11,  
wherein said electroplating step of said method comprises using an electroplating apparatus, and

wherein said electroplating apparatus comprises:

- (a) a cathode-wafer;
- (b) an anode;
- (c) an electroplating vessel; and
- (d) a voltage source.

**Claim 16 (currently amended):** A device, as recited in Claim 15,  
wherein the cathode-wafer comprises the Cu surface, and  
wherein the anode comprises at least one material selected from a group consisting essentially of copper (Cu), a copper-platinum alloy (Cu-Pt), titanium (Ti), platinum (Pt), a titanium-platinum alloy (Ti-Pt), an anodized copper-zinc alloy (Cu-Zn, i.e., brass), a

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platinized titanium (Pt/Ti), and a platinized copper-zinc (Pt/Cu-Zn, i.e., platinized brass).

**Claim 17 (canceled).**

**Claim 18 (currently amended):** A device, as recited in Claim 15, wherein said electroplating comprises a plating condition selected from a group consisting essentially of a direct voltage in the range of approximately 1 V to approximately 4 V and a direct current in the range of approximately 0.01 A to approximately 0.2 A.

**Claim 19 (currently amended):** A device, as recited in Claim 15, wherein the Zn-doping (i.e., Zn content) in the reduced-oxygen Cu-Zn alloy thin film is controllable by varying at least one electroplating condition selected from a group consisting essentially of:

increasing the at least one zinc (Zn) ion source concentration, thereby slowly increasing said Zn-doping;

increasing the at least one copper (Cu) ion source concentration, thereby slowly decreasing said Zn-doping;

increasing the solution flow rate increases Zn-doping, thereby increasing the pH decreases cathodic efficiency with respect to Zn, and thereby decreasing said Zn-doping;

increasing the electroplating duration, thereby slowly decreasing said Zn-doping;

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using a Cu anode, thereby decreasing said Zn-doping;

using a brass anode, thereby increasing said Zn-doping;

increasing the voltage, thereby increasing the Zn-doping; and

increasing the current, thereby increasing the Zn-doping.

**Claim 20 (currently amended):** A semiconductor device, having a reduced-oxygen copper-zinc alloy (Cu-Zn) thin film formed on a copper (Cu) surface, comprising:

a semiconductor substrate having at least one Cu surface formed thereon; and

a reduced-oxygen Cu-Zn alloy thin film formed, by electroplating, and disposed on the at least one Cu surface, wherein the reduced-oxygen Cu-Zn alloy thin film completely covers the at least one Cu surface,

wherein the reduced-oxygen Cu-Zn alloy thin film is formed by annealing a Cu-Zn alloy thin film in a temperature range of approximately 150°C to approximately 450°C, and

wherein the reduced-oxygen Cu-Zn alloy thin film is formed by annealing a Cu-Zn alloy thin film for a duration range of approximately 0.5 minutes to approximately 60 minutes.